Application No.: 10/735,514 2 Docket No.: 03939/100M269-US1

## **AMENDMENTS TO THE CLAIMS**

1. (Original) A method of treating movement disorders comprising administering to a subject in need of treatment, a composition comprising an amount effective for this purpose of a compound according to the following formula:

$$R_1$$
 $R_2$ 
 $R_3$ 
 $R_4$ 

or a pharmaceutically acceptable salt, prodrug or metabolite thereof, wherein  $R_3$  and  $R_4$  are each independently selected from the group consisting of lower alkyl, phenyl and lower alkyl substituted phenyl, and  $R_1$  and  $R_2$  are each independently either a hydrogen atom or a radical of the formula

wherein R<sub>5</sub> and R<sub>6</sub> are each independently selected from the group consisting of H, lower alkyl, phenyl and lower alkyl substituted phenyl.

2. (Original) The method according to claim 1, wherein R<sub>5</sub> and R<sub>6</sub> are each independently H or lower alkyl.

3. (Original) The method according to claim 1, wherein  $R_3$  and  $R_4$  are each independently phenyl.

4. (Original) The method according to claim 1, wherein at least one of  $R_1$  and  $R_2$  is defined by the formula

$$R_6$$
 $R_5$ 

wherein R<sub>5</sub> and R<sub>6</sub> are each independently either H or lower alkyl.

- (Original) The method according to claim 1, wherein at least one of R<sub>1</sub> and R<sub>2</sub> is
   -CH<sub>2</sub>OCH<sub>3</sub>.
  - 6. (Original) The method according to claim 1, wherein both  $R_5$  and  $R_6$  are phenyl,  $R_1$  is  $CH_2OCH_3$  and  $R_2$  is H.
  - 7. (Original) The method according to claim 1, wherein both  $R_5$  and  $R_6$  are phenyl and both  $R_1$  and  $R_2$  are  $-CH_2OCH_3$ .
  - 8. (Original) The method according to claim 1, wherein said movement disorder is essential tremor.

9. (Original) The method according to claim 1, wherein said movement disorder is Parkinson's disease.

- 10. (Original) The method of claim 1, wherein said movement disorder is a focal dystonia.
- 11. (Original) The method of claim 10, wherein said focal dystonia is writer's cramp.
- 12. (Original) The method according to claim 1, wherein said therapeutically effective amount is from between about 150 mg to about 1500 mg, administered in one or in two divided daily doses.
- 13. (Original) The method according to claim 12, wherein said therapeutically effective amount is from between about 200 mg to 1200 mg, administered in one or in two divided daily doses.
- 14. (Original) A method of treating movement disorders comprising administering to a subject in need of treatment a pharmaceutical dosage form comprising:
- a) a therapeutically effective amount of a compound according to the following formula:

$$R_1$$
 $R_2$ 
 $R_3$ 
 $R_4$ 

or a pharmaceutically acceptable salt, prodrug or metabolite thereof, wherein  $R_3$  and  $R_4$  are each independently selected from the group consisting of lower alkyl, phenyl and lower alkyl substituted phenyl, and  $R_1$  and  $R_2$  are each independently either a hydrogen atom or a radical of the formula

wherein R<sub>5</sub> and R<sub>6</sub> are each independently selected from the group consisting of H, lower alkyl, phenyl and lower alkyl substituted phenyl; and

- b) a pharmaceutically acceptable carrier.
- 15. (Original) The method according to claim 14, wherein R<sub>5</sub> and R<sub>6</sub> are each independently H or lower alkyl.
- 16. (Original) The method according to claim 14, wherein R<sub>3</sub> and R<sub>4</sub> are both phenyl.
- 17. (Original) The method according to claim 14, wherein at least one of  $R_1$  and  $R_2$  is defined by the formula

wherein  $R_5$  and  $R_6$  are each independently selected from H or lower alkyl.

Docket No.: 03939/100M269-US1

- 18. (Original) The method according to claim 14, wherein at least one of R<sub>1</sub> and R<sub>2</sub> is -CH<sub>2</sub>OCH<sub>3</sub>.
  - 19. (Original) The method according to claim 14, wherein both  $R_5$  and  $R_6$  are phenyl,  $R_1$  is  $CH_2OCH_3$  and  $R_2$  is H.
  - 20. (Original) The method according to claim 14, wherein both  $R_5$  and  $R_6$  are phenyl and both  $R_1$  and  $R_2$  are  $-CH_2OCH_3$ .
  - 21. (Original) The method according to claim 14, wherein said movement disorder is essential tremor.
  - 22. (Original) The method according to claim 14, wherein said movement disorder is Parkinson's disease.
  - 23. (Original) The method of claim 14, wherein said movement disorder is a focal dystonia.
  - 24. (Original) The method of claim 23, wherein said focal dystonia is writer's cramp.
  - 25. (Original) The method according to claim 14, wherein said therapeutically effective amount is from between about 150 mg to about 1500 mg, administered in one or two divided daily doses.

Application No.: 10/735,514 7 Docket No.: 03939/100M269-US1

26. (Original) The method according to claim 25, wherein said therapeutically effective amount is from between about 200 mg to 1200 mg, administered in one or two divided daily doses.

- 27. (Original) The method according to claim 14, wherein said dosage form is selected from the group consisting of oral, rectal, topical, sub-lingual, mucosal, nasal, ophthalmic, subcutaneous, intramuscular, intravenous, transdermal, spinal, intrathecal, intra-articular, intra-arterial, sub-arachinoid, bronchial, lymphatic, and intra-uterillean administered dosage forms.
- 28. (Original) The method according to claim 14, wherein said dosage form is an orally administered dosage form selected from the group consisting of tablet, capsule, caplet, gelcap, and syrup.
- 29. (Original) A method of treating essential tremor comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

$$R_1$$
 $R_2$ 
 $R_3$ 
 $R_4$ 

or a pharmaceutically acceptable salt, prodrug or metabolite thereof,

wherein  $R_3$  and  $R_4$  are each independently selected from the group consisting of lower alkyl, phenyl and lower alkyl substituted phenyl, and  $R_1$  and  $R_2$  are each independently either a hydrogen atom or a radical of the formula

wherein R<sub>5</sub> and R<sub>6</sub> are each independently selected from the group consisting of H, lower alkyl, phenyl and lower alkyl substituted phenyl.

- 30. (Original) The method according to claim 29, wherein  $R_5$  and  $R_6$  are each independently H or lower alkyl.
- 31. (Original) The method according to claim 29, wherein R<sub>3</sub> and R<sub>4</sub> are each independently phenyl.
- 32. (Original) The method according to claim 29, wherein at least one of  $R_1$  and  $R_2$  is defined by the formula

$$R_6$$

wherein  $R_5$  and  $R_6$  are each independently either H or lower alkyl.

33. (Original) The method according to claim 29, wherein at least one of  $R_1$  and  $R_2$  is  $-CH_2OCH_3$ .

- 34. (Original) The method according to claim 29, wherein both  $R_5$  and  $R_6$  are phenyl,  $R_1$  is  $CH_2OCH_3$  and  $R_2$  is H.
- 35. (Original) The method according to claim 29, wherein both  $R_5$  and  $R_6$  are phenyl and both  $R_1$  and  $R_2$  are  $-CH_2OCH_3$ .
- 36. (Original) The method according to claim 29, wherein said therapeutically effective amount is from between about 150 mg to about 1500 mg, administered in or two daily doses.
- 37. (Original) The method according to claim 36, wherein said therapeutically effective amount is from between about 200 mg to 1200 mg, administered in one or two daily doses.
- 38. (Original) A method of treating essential tremor comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

39. (Original) A method of treating essential tremor comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

40. (Original) A method of treating essential tremor comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

41. (Original) A method of treating Parkinson's disease comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

$$R_1$$
 $R_2$ 
 $R_3$ 
 $R_4$ 

or a pharmaceutically acceptable salt, prodrug or metabolite thereof,

Application No.: 10/735,514

12 I

wherein  $R_3$  and  $R_4$  are each independently selected from the group consisting of lower alkyl, phenyl and lower alkyl substituted phenyl, and  $R_1$  and  $R_2$  are each independently either a hydrogen atom or a radical of the formula

wherein R<sub>5</sub> and R<sub>6</sub> are each independently selected from the group consisting of H, lower alkyl, phenyl and lower alkyl substituted phenyl.

- 42. (Original) The method according to claim 41, wherein R<sub>5</sub> and R<sub>6</sub> are each independently H or lower alkyl.
- 43. (Original) The method according to claim 41, wherein R<sub>3</sub> and R<sub>4</sub> are each independently phenyl.
- 44. (Original) The method according to claim 41, wherein at least one of  $R_1$  and  $R_2$  is defined by the formula

wherein  $R_5$  and  $R_6$  are each independently either H or lower alkyl.

Docket No.: 03939/100M269-US1

- 46. (Original) The method according to claim 41, wherein both  $R_5$  and  $R_6$  are phenyl,  $R_1$  is  $CH_2OCH_3$  and  $R_2$  is H.
- 47. (Original) The method according to claim 41, wherein both  $R_5$  and  $R_6$  are phenyl and both  $R_1$  and  $R_2$  are  $-CH_2OCH_3$ .
- 48. (Original) The method according to claim 41, wherein said therapeutically effective amount is from between about 150 mg to about 1500 mg, administered in or two daily doses.
- 49. (Original) The method according to claim 48, wherein said therapeutically effective amount is from between about 200 mg to 1200 mg, administered in or two daily doses.
- 50. (Original) A method of treating Parkinson's disease comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

51. (Original) A method of treating Parkinson's disease comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

Application No.: 10/735,514

52. (Original) A method of treating Parkinson's disease comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

15

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

- 53. (Original) The method of claim 1 wherein said movement disorder is selected from the group consisting of tremor, dystonia, chorea, athetosis, a tic disorder, blepharospasm, hemiballysmus, myoclonus, torticollis, writer's cramp, restless leg syndrome and asterixis.
- 54. (Orginal) The method of claim 1 wherein said movement disorder is selected from the group consisting of Parkinson's disease, Tourette's syndrome, progressive supranuclear palsy and Wilson's disease.
- 55. (Original) A method of treating movement disorders comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

- 56. (Original) The method of claim 55, wherein the movement disorder is essential tremor.
- 57. (Original) A method of treating movement disorders comprising administering to a subject in need of treatment a composition comprising a therapeutically effective amount of a compound according to the following formula:

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

58. (Original) The method of claim 57, wherein the movement disorder is essential tremor.

Docket No.: 03939/100M269-US1

Application No.: 10/735,514

59. (Currently Amended) A method of treating essential tremor movement disorders comprising administering to a subject in need of treatment a composition comprising a therapeutically

17

or a pharmaceutically acceptable salt, prodrug or metabolite thereof.

effective amount of a compound according to the following formula:

60. (Original) The method of claim 59, wherein the movement disorder is essential tremor.